

Notice of References Cited

Application/Control No.

10/021,955

Applicant(s)/Patent Under
Reexamination
LUPSKI ET AL.

Examiner

Suryaprabha Chunduru

Art Unit

1637

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*		Document Number Country Code-Number-Kind Code	Date MM-YYYY	Name	Classification
	A	US-5,691,144	11-1997	Boss et al.	435/6
*	B	US-5,780,223	07-1998	Lupski et al.	435/6
	C	US-			
	D	US-			
	E	US-			
	F	US-			
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	I	US-			
	J	US-			
	K	US-			
	L	US-			
	M	US-			

FOREIGN PATENT DOCUMENTS

*		Document Number Country Code-Number-Kind Code	Date MM-YYYY	Country	Name	Classification
	N					
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NON-PATENT DOCUMENTS

*		Include as applicable: Author, Title Date, Publisher, Edition or Volume, Pertinent Pages)
*	U	Timmerman et al. Novel missense mutation in the early growth response 2 gene associated with Dejerine-Sttas syndrome phenotype. Neurology, Vol. 52, pp. 1827-1832, 1999.
*	V	Gillespie et al. Periaxin, a novel protein of myelinating schwann cells with a possible role in axonal ensheathment. Neuron, Vol 12, pp. 497-508, 1994.
	W	Guibot et al. A mutation in periaxin is responsible for CNT4F, an autosomal recessive form of Charcot-Marie-Tooth disease. Human Mol Genetics, Vol. 10, No. 4, pp. 415-421, 2001.
*	X	Roa et al. Dejerine-Scottas syndrome associated with point mutation in the peripheral myelin protein 22 (PMP22) gene. Nature Genetics, Vol. 5, pp. 269-273, 1993.

*A copy of this reference is not being furnished with this Office action. (See MPEP § 707.05(a).)
Dates in MM-YYYY format are publication dates. Classifications may be US or foreign.